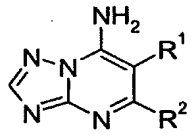
**Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in the application:

**Listing of Claims:**

1. (Currently amended) A triazolopyrimidine of the formula I



I

in which the substituents are as defined below:

$R^1$ is $C_5$-$C_{12}$-alkyl or $C_5$-$C_{14}$-alkoxyalkyl, where the aliphatic groups may be substituted by 1 to 3 of the following groups:

cyano, nitro, hydroxyl, $C_3$-$C_6$-cycloalkyl, $C_1$-$C_6$-alkylthio, and $NR^aR^b$;

$R^a$, $R^b$are hydrogen or $C_1$-$C_6$-alkyl;

$R^2$ is $CHR^3CH_3$, ~~cyclopropyl, $CH=CH_2$~~ or $CH_2CH=CH_2$;
$R^3$ is hydrogen or $CH_3$.

2. (Original) The compound of the formula I according to claim 1, in which $R^2$ is ethyl.

3. (Previously presented) The compound of the formula I according to claim 1, in which $R^1$ is an unsubstituted straight-chain or mono-, di- or tribranched alkyl chain having up to 12 carbon atoms.

4. (Previously presented) Triazolopyrimidines of the formula I according to claim 1 selected from the group consisting of:
5-ethyl-6-(1-methylheptyl)-[1 ,2,4]triazolo[1 ,5-a]pyrimidin-7-ylamine;
5-ethyl-6-octyl-[1 ,2,4]triazolo[1 ,5-a]pyrimidin-7-ylamine;
5-ethyl-6-(3,5,5-trimethylhexyl)-[1 ,2,4]triazolo[1 ,5-a]pyrimidin-7-ylamine;

5-ethyl-6-pentyl-[1 ,2,4]triazolo[1 ,5-a]pyrimidin-7-ylamine;

5-ethyl-6-hexyl-[1 ,2,4]triazolo[1 ,5-a]pyrimidin-7-ylamine;

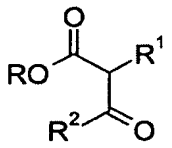
5-ethyl-6-heptyl-[1 ,2,4]triazolo[1 ,5-a]pyrimidin-7-ylamine;

5-ethyl-6-nonyl-[1 ,2,4]triazolo[1 ,5-a]pyrimidin-7-ylamine;

5-ethyl-6-undecyl-[1 ,2,4]triazolo[1 ,5-a]pyrimidin-7-ylamine;

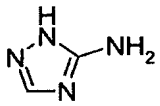
5-ethyl-6-(3-pentyloxypropyl)-[1,2,4]triazolo[1,5-a]pyrimidin-7-ylamine.

5. (Previously presented) A process for preparing compounds of the formula I according to any of claims 1 to 4 wherein B-keto esters of the formula II,



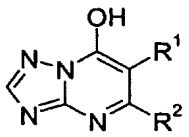
II

in which R is $C_1$-$C_4$-alkyl are reacted with 3-amino-1 ,2,4-triazole of the formula III



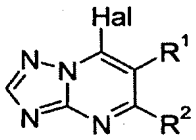
III

to give 7-hydroxytriazolopyrimidines of the formula IV



IV

which are halogenated to give compounds of the formula V

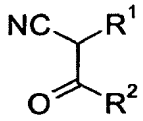


V

in which Hal is chlorine or bromine and V is reacted with ammonia.

6. (Previously presented) A compound of the formula IV or V according to claim 5.

7. (Previously presented) A process for preparing compounds of the formula I according to any of claims 1 to 4 wherein acyl cyanides of the formula VI,



VI

are reacted with 3-amino-1 ,2,4-triazole of the formula III.

8. (Previously presented) A fungicidal composition comprising a solid or liquid carrier and a compound of the formula I according to any of claims 1 to 4.

9. (Previously presented) Seed comprising a compound of the formula I according to any of claims 1 to 4 in an amount of 1 to 1000 g per 100 kg.

10. (Previously presented) A method for controlling phytopathogenic harmful fungi, wherein the fungi or the materials, plants, the soil or seed to be protected against fungal attack are treated with an effective amount of the compound of the formula I according to any of claims 1 to 4.

11. (Canceled)

12. (Canceled)

13. (Canceled)